

510(k) Summary for the 3S Hemi Toe Implant

In accordance with 21 CFR 807.92 of the Federal Code of Regulations
the following 510(k) summary is submitted for the 3S Hemi Toe Implant.

Date Prepared: October 11, 2007

1. Submitter:

Trilliant Surgical LTD
1630 W. 13th Street03
Houston, TX 77008

Contact Person:

J.D. Webb
The OrthoMedix Group, Inc.
1001 Oakwood Blvd
Round Rock, TX 78681
Telephone: 512-388-0199

NOV 28 2007

2. Trade name: 3S Hemi Toe Implant

Common Name: Hemi toe

Classification Name: prosthesis, toe, hemi-, phalangeal
Class II per 21 CFR section 888.3730
KWD

3. Predicate or legally marketed devices which are substantially equivalent:

Futura Biomedical Metal Hemi Toe Implant (K971047), Kinetikos Medical K2 Hemi Toe Implant System (K023770), and Townley Great Toe Joint (K911378).

4. Description of the device:

The 3S Hemi Toe is a single stemmed resurfacing prosthesis for the first proximal phalanx designed to supplement first metatarsal phalangeal joint arthroplasty. The concave congruent articular surface has a mirror finish to minimize friction and matches the adjacent metatarsal head. The oval shape helps to reduce impingement on the metatarsal head, maintain range of motion and reduce pain without altering the joint biomechanics. The 3S Hemi Toe requires minimal bone resection and provides full range of motion of the first metatarsophalangeal joint (MPJ).

Materials:

The substrate of the 3S Hemi Toe is fabricated from CoCrMo alloy per ASTM F799 or F75.

5. Intended Use:

The 3S Hemi Toe Implant is designed to supplement first metatarsal phalangeal joint arthroplasty. Indications include hallux limitus or hallux rigidus, painful hallux valgus, revision of failed previous surgery and painful arthritis. This device is for uncemented use.

6. Comparison of the technological characteristics of the device to predicate and legally marketed devices:

The 3S Hemi Toe does not incorporate any new technological characteristics as compared to the predicate device. The 3S Hemi Toe and the predicate devices are made from the same material.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

NOV 28 2007

Trilliant Surgical Ltd.
% Mr. J. D. Webb
The OrthoMedix Group, Inc.
1001 Oakwood Blvd.
Round Rock, Texas 78681

Re: K072922

Trade/Device Name: 3S Hemi Toe Implant
Regulation Number: 21 CFR 888.3730
Regulation Name: Toe joint phalangeal (hemi-toe) polymer prosthesis
Regulatory Class: II
Product Code: KWD
Dated: October 11, 2007
Received: October 15, 2007

Dear Mr. Webb:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set

forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050. This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Center for Devices and Radiological Health's (CDRH's) Office of Compliance at (240) 276-0120. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding postmarket surveillance, please contact CDRH's Office of Surveillance and Biometric's (OSB's) Division of Postmarket Surveillance at (240) 276-3474. For questions regarding the reporting of device adverse events (Medical Device Reporting (MDR)), please contact the Division of Surveillance Systems at (240) 276-3464. You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (240) 276-3150 or at its Internet address <http://www.fda.gov/cdrh/industry/support/index.html>.

Sincerely yours,



Mark N. Melkerson
Director
Division of General, Restorative
and Neurological Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

Indications for Use

510(k) Number (if known): K072922

Device Name: 3S Hemi Toe Implant

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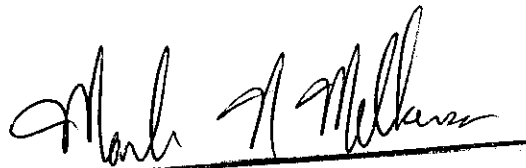
Prescription Use X
(Part 21 CFR 801 Subpart D)

AND/OR

Over-The-Counter Use _____
(21 CFR 807 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE
OF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)



Division Sign-Off

Division of General, Restorative,
and Neurological Devices

510(k) Number K072922